Clearance Officer, United States Department of Justice, Information Management and Security Staff, Justice Management Division, Suite 1220, Washington Center, 1331 G Street, NW, Washington, DC 20530.

Dated: December 21, 1999.

Brenda E. Dyer,

Department Deputy Clearance Officer, Department of Justice.

[FR Doc. 99-33549 Filed 12-27-99; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application

Pursuant to Section 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on November 4, 1999, Celgene Corporation, 7 Powder Horn Drive, Warren, New Jersey 07059, made application by renewal to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of methylphenidate (1724) a basic class of controlled substance listed in Schedule II.

The firm plans to manufacture methylphenidate for product research and development.

Any other such applicant and any person who is presently registered with DEA to manufacture such substance may file comments or objections to the issuance of the proposed registration.

Any such comments or objections may be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, DC 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than February 28, 2000.

Dated: December 16, 1999.

John H. King,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 99–33648 Filed 12–27–99; 8:45 am] BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated August 6, 1999, and published in the **Federal Register** on August 20, 1999, (64 FR 45564), Guilford Pharmaceuticals, Inc., 6611 Tributary Street, Baltimore, Maryland 21224, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of cocaine (9041), a basic class of controlled substance listed in Schedule II

The firm plans to manufacture methyl-3-beta-(4-trimethylstannylphenyl)-tropane-2-carboxylate as a final intermediate for the production of dopascan injection.

DEA has considered the factors in Title 21, United States Code, Section 823(a) and determined that the registration of Guilford Pharmaceuticals to manufacture the listed controlled substance is consistent with the public interest at this time. DEA has investigated the firm on a regular basis to ensure that the company's continued registration is consistent with the public interest. These investigations have included inspection and testing of the conpany's physical security systems, audits of the company's records, verification of the company's compliance with state and local laws, and a review of the company's background and history. Therefore, pursuant to 21 U.S.C. 823 and 28 CFR 0.100 and 0.104, the Deputy Assistant Administrator, Office of Diversion Control, hereby orders that the application submitted by the above firm for registration as a bulk manufacturer of the basic class of controlled substance listed above is granted.

Dated: December 16, 1999.

John H. King,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 99–33645 Filed 12–27–99; 8:45 am] $\tt BILLING\ CODE\ 4410-09-M$

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. 96-10]

Wesley G. Harline, M.D.; Continuation of Registration With Restrictions

On October 27, 1995, the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration (DEA) issued an Order to Show Cause to Wesley Harline, M.D. (Respondent) of Ogden, Utah, notifying him of an opportunity to show cause as to why DEA should not revoke his DEA Certificate of Registration AH1650248 and deny any pending applications for renewal of such registration as a practitioner pursuant to 21 U.S.C. 823(f) and 824(a)(4), for reason that his

continued registration would be inconsistent with the public interest.

By letter dated December 14, 1995, Respondent, through counsel, filed a request for a hearing, and following prehearing procedures, a hearing was held in Salt Lake City, Utah on April 1 through 3 and May 6 through 8, 1997, and by telephone in Salt Lake City and Arlington, Virginia, on August 18 through 21, 1997, before Administrative Law Judge Mary Ellen Bittner. At the hearing both parties called witnesses to testify and introduced documentary evidence. After the hearing both parties submitted proposed findings of fact, conclusions of law and argument.

In this brief, Respondent's counsel included findings based upon evidence that was not introduced at the hearing. On January 5, 1998, the Government filed a Motion to Strike Post Record Evidence from Respondent's Proposed Findings of Fact, Conclusions of Law and Argument. On January 21, 1998, Respondent filed his Opposition to Government's Motion to Strike Post Record Evidence, and in the alternative, Motion to Reopen the Record.

On April 2, 1999, Judge Bittner issued her Opinion and Recommended Ruling Findings of Fact, Conclusions of Law and Decision (Opinion), granting the Government's motion to strike the additional evidence, denving Respondent's motion to reopen the record, and recommending that Respondent's DEA Certificate of Registration be revoked and any pending applications be denied. On June 14, 1999, Respondent filed exceptions to Judge Bittner's Opinion and on August 2, 1999, the Government filed its response to Respondent's exceptions. Thereafter, on August 10, 1999, Judge Bittner transmitted the record of these proceedings to the Deputy Administrator.

While this matter was pending with the Deputy Administrator, Respondent submitted a letter dated November 4, 1999, responding to the Government's response to his exceptions and formally moving that the record be reopened to allow additional evidence to be considered. As will be discussed more fully below, the Acting Deputy Administrator denies Respondent's motion to reopen the record and has not considered Respondent's letter dated November 4, 1999, in rendering his decision in this matter.

The Acting Deputy Administrator has considered the record in its entirety, and pursuant to 21 CFR 1316.67, hereby issues his final order based upon findings of fact and conclusions of law as hereinafter set forth. The Acting Deputy Administrator adopts, except as